

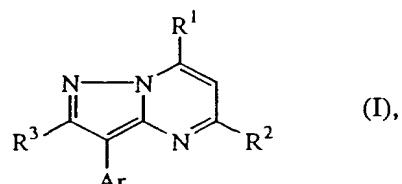
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ABSTRACT

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PYRAZOLOPYRIMIDINES  
AS CRF RECEPTOR ANTAGONISTS

This invention concerns compounds of formula



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including the stereoisomers and the pharmaceutically acceptable acid addition salt forms thereof, wherein R<sup>1</sup> is NR<sup>4</sup>R<sup>5</sup> or OR<sup>5</sup>; R<sup>2</sup> is C<sub>1</sub>-6alkyl, C<sub>1</sub>-6alkyloxy or C<sub>1</sub>-6alkylthio; R<sup>3</sup> is hydrogen, C<sub>1</sub>-6alkyl, C<sub>1</sub>-6alkylsulfonyl, C<sub>1</sub>-6alkylsulfoxyl or C<sub>1</sub>-6alkylthio; R<sup>4</sup> is hydrogen, C<sub>1</sub>-6alkyl, mono- or di(C<sub>3</sub>-6cycloalkyl)methyl,  
15 C<sub>3</sub>-6cycloalkyl, C<sub>3</sub>-6alkenyl, hydroxyC<sub>1</sub>-6alkyl, C<sub>1</sub>-6alkylcarbonyloxyC<sub>1</sub>-6alkyl or C<sub>1</sub>-6alkyloxyC<sub>1</sub>-6alkyl; R<sup>5</sup> is C<sub>1</sub>-8alkyl, mono- or di(C<sub>3</sub>-6cycloalkyl)methyl, Ar<sup>1</sup>CH<sub>2</sub>, C<sub>1</sub>-6alkyloxyC<sub>1</sub>-6alkyl, hydroxyC<sub>1</sub>-6alkyl, C<sub>3</sub>-6alkenyl, thienylmethyl, furanylmethyl, C<sub>1</sub>-6alkylthioC<sub>1</sub>-6alkyl, morpholinyl, mono- or di(C<sub>1</sub>-6alkyl)aminoC<sub>1</sub>-6alkyl, di(C<sub>1</sub>-6alkyl)amino, C<sub>1</sub>-6alkylcarbonylC<sub>1</sub>-6alkyl, C<sub>1</sub>-6alkyl substituted with imidazolyl;  
20 or a radical of formula -Alk-O-CO-Ar<sup>1</sup>; or R<sup>4</sup> and R<sup>5</sup> taken together with the nitrogen atom to which they are attached may form an optionally substituted pyrrolidinyl, piperidinyl, homopiperidinyl or morpholinyl group; having CRF receptor antagonistic properties; pharmaceutical compositions containing such compounds as active ingredients; methods of treating disorders related to hypersecretion of CRF such as  
25 depression, anxiety, substance abuse, by administering an effective amount of a compound of formula (I).